

**K232468 SC Medica FFX**May 9, 2024  
268 days to decisionK232468 · Product code: **MRW** · Orthopedic  
Source: <https://www.510kdatabase.net/k232468/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	Aug 15, 2023
Decision date	May 9, 2024
Days to decision	268 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sc Medica</b>
Location	Strasbourg, FR
Contact	Camille Srour
510(k) history	3 submissions · 3 cleared · 2024-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k232468/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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